

CLAIMS

Sub A1

1. A composite intended for medical use, in particular surgical or therapeutic use, characterized in that it comprises

5 - a thermoplastic component plasticizable within the temperature range -10 °C...+100 °C, which is substantially made up of hydroxy acids and structural units derived from hydroxy acid derivatives, and the molar mass of which is within the range 10,000 - 1,000,000 g/mol, and which degrades in the body typically within a period ranging from 10 a few days to several years, and which in its solid state is a mechanically strong plastic or rubbery material, and - a bioactive component, which is a bioactive glass, a bioactive xerogel, a bioactive ceramic material, coral or a coral-based product, or a bioactive glass ceramic material.

15 2. The composite according to Claim 1, characterized in that the plastic component is plasticizable within the temperature range 5 °C...70 °C, preferably within the temperature range 37 °C...55 °C.

20 3. The composite according to Claim 1 or 2, characterized in that the plasticized plastic component remains moldable for a certain period even after the temperature of the composite has been lowered to a temperature which is considerably lower than the setting temperature of the said plastic component.

25 34. The composite according to Claim 1, 2 or 3, characterized in that the plastic component is biodegradable in a controlled manner within the time range 1 week - 3 years.

30 45. The composite according to Claim 3⁴, characterized in that the structural unit is an L-, D- or DL-lactic acid; an L-, D- or DL-lactide; or epsilon-caprolactone.

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5. The composite according to Claim 5, characterized in that the plastic component is a copolymer based on structural units of L-lactide and epsilon-caprolactone.

6. The composite according to Claim 5, characterized in that the composition of the copolymer is within the range

10240
epsilon-caprolactone
----- = 2/98 ... 98/2
L-lactide

7. The composite according to Claim 7, characterized in that

10241
epsilon-caprolactone
----- = 4 : 1
L-lactide

8. The composite according to Claim 8, characterized in that the molar mass of the copolymer is approx. 30,000 - 300,000 g/mol.

9. The composite according to *Claim 1*, characterized in that the bioactive component is present as separate particles in the composite.

10. The composite according to *Claim 10*, characterized in that the separate particles are fibers, porous pieces, microparticles or glass beads.

11. The composite according to *Claim 1*, characterized in that the plastic component and/or the bioactive component contains one or more additives.

12. The composite according to *Claim 1*, characterized in that the plastic component and the bioactive component form a dense piece.

13. The composite according to *Claim 1*, characterized in that the plastic component and the bioactive component form a dense piece.

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characterized in that the plastic component forms a porous piece.

14/14 16. A blend intended for the preparation of a composite according to ^{Claim 1} any of Claims 1 - 14, characterized in that the 5 plastic component and the bioactive component in the blend are in powder form.

15/15 16. A coating, membrane, net, powder, fiber, thread, adhesive, or a piece such as a plate, bead, tube, nail or rod, prepared from the composite according to ^{Claim 1} any of Claims 10 1 - 14.

17. The use of a composite according to ^{Claim 1} any of Claims 1 - 14 for the preparation of any of the following products:

- a bone or cartilage application, such as a filling material for bone or cartilage, a product intended for the repairing of long bones, a plate for the repairing of the back of the eye or facial bones, a bone cement, an adhesive for joining the product to a tissue or tissues, an implant coating, a piece for the repairing of the vertebral column, and a skull plate,
- a tooth or jaw application, such as a temporary tooth filling material, a temporary or permanent tooth root filling material, a parodontal product, a product to be placed in the cavity left by an extracted tooth, a tooth cement, a temporary tooth cement, a temporary crown material, a tooth implant coating, an occlusion index rail, a surgical paste, and a template material, which may be, for example, a paste, ring or thread to be fitted in a gingival pocket,
- a cartilage coating,
- a tissue guiding membrane or tube,
- a protective cloth, a wound dressing, or an adhesive tape,
- a carrier for an active agent, such as a drug, or for some other biological structure.

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